



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

WT1 expression in breast cancer disrupts the epithelial/mesenchymal balance of tumour cells and correlates with the metabolic response to docetaxel

Citation for published version:

Artibani, M, Sims, AH, Slight, J, Aitken, S, Thornburn, A, Muir, M, Brunton, VG, Del-Pozo, J, Morrison, LR, Katz, E, Hastie, ND & Hohenstein, P 2017, 'WT1 expression in breast cancer disrupts the epithelial/mesenchymal balance of tumour cells and correlates with the metabolic response to docetaxel', *Scientific Reports*, vol. 7, 45255. <https://doi.org/10.1038/srep45255>

Digital Object Identifier (DOI):

[10.1038/srep45255](https://doi.org/10.1038/srep45255)

Link:

[Link to publication record in Edinburgh Research Explorer](#)

Document Version:

Publisher's PDF, also known as Version of record

Published In:

Scientific Reports

Publisher Rights Statement:

© The Author(s) 2017. This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>

General rights

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.



SCIENTIFIC REPORTS

OPEN

***WT1* expression in breast cancer disrupts the epithelial/mesenchymal balance of tumour cells and correlates with the metabolic response to docetaxel**

Mara Artibani^{1,†}, Andrew H. Sims¹, Joan Slight¹, Stuart Aitken¹, Anna Thornburn¹, Morwenna Muir¹, Valerie G. Brunton¹, Jorge Del-Pozo², Linda R. Morrison², Elad Katz^{1,‡}, Nicholas D. Hastie¹ & Peter Hohenstein^{1,3}

WT1 is a transcription factor which regulates the epithelial-mesenchymal balance during embryonic development and, if mutated, can lead to the formation of Wilms' tumour, the most common paediatric kidney cancer. Its expression has also been reported in several adult tumour types, including breast cancer, and usually correlates with poor outcome. However, published data is inconsistent and the role of **WT1** in this malignancy remains unclear. Here we provide a complete study of **WT1** expression across different breast cancer subtypes as well as isoform specific expression analysis. Using *in vitro* cell lines, clinical samples and publicly available gene expression datasets, we demonstrate that **WT1** plays a role in regulating the epithelial-mesenchymal balance of breast cancer cells and that **WT1**-expressing tumours are mainly associated with a mesenchymal phenotype. **WT1** gene expression also correlates with **CYP3A4** levels and is associated with poorer response to taxane treatment. Our work is the first to demonstrate that the known association between **WT1** expression in breast cancer and poor prognosis is potentially due to cancer-related epithelial-to-mesenchymal transition (EMT) and poor chemotherapy response.

The Wilms' Tumour gene 1 (**WT1**), encodes for a complex protein with transcription factor activity which is essential in mammals throughout life¹. Its roles vary with the developmental stages: in the embryo, it regulates the epithelial-mesenchymal balance required for a correct organogenesis and acts as a tumour suppressor in the aetiology of Wilms' tumour^{2,3}; in the adult, it is involved in the maintenance of tissue homeostasis and has been controversially considered as an oncogene in leukaemia and several solid tumours, including breast cancer⁴. The role of **WT1** in this malignancy is unclear, despite the accumulation of a considerable body of data over the last fifteen years. Several groups have reported **WT1** expression in breast cancer, but the data on the percentage of **WT1**-positive tumours are highly discordant, most likely because of differences in the detection method, antibody specificity and histological subtype^{5–11}. Moreover, there is evidence for both oncogenic and tumour suppressing functions^{6,7,12} and these contradictory results can be explained only in part by the presence of different isoforms¹³.

One study has associated **WT1** with improved outcomes¹⁴, however most literature seems to indicate that high **WT1** expression is associated with poor prognosis in breast cancer patients^{8,15}. The biological basis behind these seemingly inconsistent findings has not yet been identified, but several hypotheses have been formulated. Firstly, tumours expressing high levels of **WT1** may have a proliferative advantage since silencing **WT1** leads to growth

¹Institute of Genetics and Molecular Medicine, University of Edinburgh, Crewe Road South, Edinburgh, EH4 2XU, UK.

²Royal (Dick) School of Veterinary Studies, University of Edinburgh, Easter Bush Campus, Midlothian, EH25 9RG, UK.

³The Roslin Institute, University of Edinburgh, Easter Bush Campus, Midlothian, EH25 9RG, UK. [†]Present address: Weatherall Institute of Molecular Medicine, University of Oxford, OX3 9DS, UK. [‡]Present address: Drug Discovery Unit, School of Life Sciences, University of Dundee, Dow Street, Dundee DD1 5EH, UK. Correspondence and requests for materials should be addressed to P.H. (email: peter.hohenstein@roslin.ed.ac.uk)

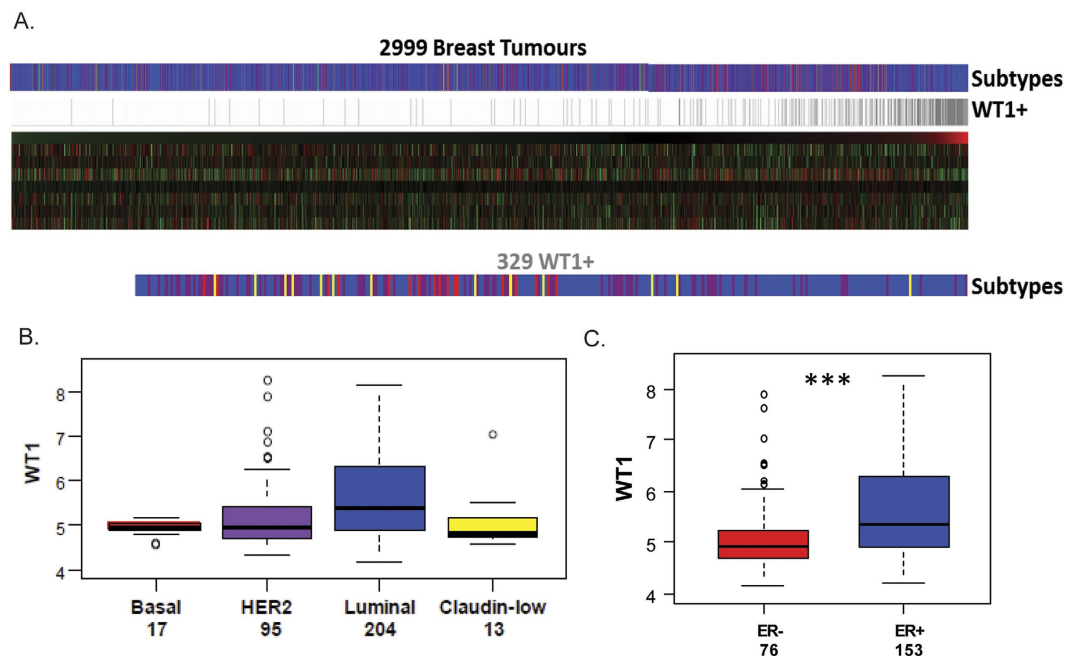


Figure 1. *WT1* expression in human primary breast cancer datasets. (A) Gene expression analysis data from 17 integrated datasets ($n = 2999$ tumours). (B) Boxplot of *WT1* expression in the different subtypes, y-axis showing log₂ values. (C) Boxplots of *WT1* expression in ER-positive (blue) and ER-negative (red) tumours designated by IHC of ER-alpha amongst those tumours with detectable expression of *WT1*, *** $p < 0.001$.

arrest and apoptosis^{16–19}. Investigations on the molecular mechanisms involved in this process have revealed that *WT1* can modulate many genes of the *BCL2* family, including *BAK*, *BAG3*, *BCL2A1* and *BCL2* itself^{20–22} as well as regulate the Fas-related death signalling pathway²³; moreover, there is some evidence suggesting that *WT1* can promote cell proliferation by up-regulating cyclin D1²⁴.

A second hypothesis is related to blood vessel formation. *WT1* has been detected in the vasculature of different tumour types^{25,26} and its expression in endometrial cancer has been associated with the induction of angiogenesis²⁷. In addition, *WT1* directly upregulates the expression of the vascular endothelial growth factor (*VEGF*)²⁸ as well as regulating its splicing²⁹. Many other vascular genes have been identified as potential *WT1* targets, including the vascular endothelial cadherin, angiopoietin-2, vascular endothelial growth factor receptors 1 and 2³⁰. Most importantly, the endothelial knockout of *WT1* has been shown to reduce both tumour angiogenesis and growth *in vivo*³¹, suggesting that this gene is a major regulator of blood vessel formation in cancer.

Finally, an alternative hypothesis relies on the fact that *WT1* is a key regulator of the epithelial/mesenchymal balance during development and therefore it may also play a role in the EMT of tumour cells¹. Besides being recently linked to chemoresistance^{32,33}, cancer-related EMT has long been associated with the acquisition of a malignant phenotype by the epithelial tumour cells: features of EMT have been described in breast³⁴ and colorectal cancer³⁵, mainly at the invasive front of the tumour. This suggests that the EMT may generate migratory cells which leave the primary site, invade the blood vessels and potentially metastasise. This theory on EMT contribution to tumour progression is supported by the fact that many developmental EMT drivers, including *SNAIL*, *SLUG*, *TWIST* and *ZEB1*, are aberrantly expressed in cancer and significantly correlate with relapse and poor clinical outcomes^{36–38}. Importantly, *WT1* has been shown to regulate the EMT which occurs in the developing epicardium² as well as the MET which is required for nephrogenesis³ and its transcriptional targets include *SNAIL*, *SLUG* and *TGF- β* ^{32,39,40}.

In the present work, we have studied the relationship between *WT1* expression and the overall epithelial/mesenchymal balance of breast cancer using *in vitro* cell lines, clinical samples and publicly available gene expression datasets in order to better investigate why *WT1* is usually associated with poor prognosis.

Results

Different *WT1* isoforms are observed in breast cancer and their expression is higher in the ER-positive subtype. Several groups have described *WT1* expression in breast cancer^{5–11}, however the data is inconsistent and the frequency of *WT1*-positive tumours varies from 23%⁶ to 87%⁷. Such discrepancies may be due to different detection techniques or antibody specificity¹¹, but may also be related to differences in the molecular subtype of the tumours or to different *WT1* isoforms.

In order to overcome these limitations, we have performed a more detailed and comprehensive analysis of *WT1* expression in breast cancer using different approaches (*in vivo*, *in vitro* and *in silico*).

Firstly, we analysed *WT1* expression in an integrated dataset from 17 studies representing 2999 primary human breast cancers⁴¹ (Fig. 1A). The transcript could only be reliably detected (considering ‘Present’ detection calls in the Affymetrix data) in 11% of the tumours (329 out of the total 2999) and showed the highest expression

level in the luminal subtype (Fig. 1B, Mann Whitney p value of 4.5×10^{-7}). Furthermore, restricting the samples to those in which *WT1* was detected, *WT1* expression was significantly higher in ER-positive than in ER-negative tumours (Fig. 1C, $p = 1.2 \times 10^{-5}$).

Secondly, we analysed a locally derived cohort of invasive breast cancer samples ($n = 44$) using isoform-specific assays: quantitative RT-PCR was first performed using primers targeting exon 7/8 of *WT1*, in order to amplify all the known isoforms (Fig. 2A, Figure S1 in Additional file 1). Additional assays were then designed to specifically amplify a maternally-imprinted alternative exon 1⁴² and the portion of intron 5 which is transcribed in the truncated variant⁴³ (Figure S1 in Additional file 1). To compare the expression levels of the different isoforms between healthy and neoplastic tissue, we also included a sample of human adult mammary gland. This normal tissue sample, a pool from 5 different healthy donors, expressed higher *WT1* levels than any of the tumour samples, but lacked both the exon 1A and intron 5 isoforms.

Lastly, examination of *WT1* expression in a panel of thirteen breast cancer cell lines by qPCR demonstrated that seven of the cell lines had detectable levels of *WT1*, with the highest expression observed in the basal triple-negative lines HBL100 and MDA-MB-157 and in the luminal ER+ line ZR75 (Fig. 2B). While the qPCR for the optional exon 5 mirrors the exon 7/8 results, only BT20 and MDA-MB-231 cells expressed detectable levels of the intron 5 isoform and exon 1A was expressed only by BT20 cells.

To further assess the extraordinary complexity of the *WT1* locus, we also analysed a dataset representing a panel of 41 breast cancer cell lines using Affymetrix exon GeneChips (GSE16732)⁴⁴.

The exon array data was largely in agreement with the qPCR experiments, showing high levels of *WT1* expression for ZR75, MDA-MB-157, HS578T and T47D cell lines (Fig. 2C); moreover, *WT1* expression in MDA-MB-231 cells seemed to be restricted to exons 6 to 10, suggesting that this cell line may only express the truncated intron 5 isoform.

Knock-down of *WT1* in the MDA-MB-157 line increases the motility, invasiveness and metastatic potential of breast cancer cells. An RNAi-based gene knockdown strategy was used to investigate the functional role of *WT1* in breast cancer cells. Unlike previous studies based on transient transfections⁴⁵, we achieved stable *WT1* knockdown in three human breast cancer cell lines (Fig. 3A,B). Both constitutive and inducible lentiviral vectors were used for MDA-MB-157 cells, which showed up to 70% mRNA and protein reduction (Fig. 3A–D).

RNA-seq was used to investigate the consequences of *WT1* loss in the MDA-MB-157 cell line by comparing the transcriptomes of the wildtype and knockdown cells. Differential expression of 427 transcripts (Table S1 in Additional file 2) was observed, including several genes involved in RNA metabolism/biosynthesis, the neuronal gene *SEMA5A* (reported to play a role in cancer progression⁴⁶), and the ncRNA *MALAT1* (a regulator of the metastasis phenotype in lung cancer⁴⁷). ToppFun (<http://toppgene.cchmc.org/enrichment.jsp>)⁴⁸ was used to functionally annotate the differentially expressed genes and it showed enrichment for genes involved in several migration-related biological processes, from cell adhesion to cell projection morphogenesis/organisation (Table S2 in Additional file 2).

In order to assess if *WT1* knockdown affects the motility of breast cancer cells, we performed migration and invasion assays. The cells in which *WT1* was silenced displayed a statistically significant increase in both migratory capacity and invasiveness when compared to pGIPZ-lacZ controls ($p = 0.031$ and $p = 0.008$ respectively) (Fig. 4A).

The knockdown cells also showed significantly increased incidence of lung metastases following injection into the tail vein of nude athymic mice ($p = 0.011$) (Fig. 4B), although the total tumour area was not significantly different between the two groups (Fig. 4C).

Knockdown and overexpression of *WT1* disrupts the epithelial-mesenchymal balance of breast cancer cells. Since increased migration is considered a feature of EMT⁴⁹, we determined if the loss of *WT1* also affects the epithelial/mesenchymal balance of breast cancer cells and analysed the RNA levels of several EMT drivers (*SNAI1*, *SNAI2*, *ZEB1*, *ZEB2*, *TWIST1*), epithelial markers (*CTNNA1*, *CTNNA2*, *CTNND1*, *CDH1*, *KRT18*) and mesenchymal markers (*VIM*, *TNC*, *TGFB1*, *CDH2*, *MMP2*, *MMP9*) through quantitative real-time PCR.

We also generated and analysed overexpression clones from two different lines using the full length cDNA of *WT1* (Fig. 3E).

In the mesenchymal line MDA-MB-157, overexpressing *WT1* pushes the cells towards a more epithelial phenotype, with increased expression of *CTNNA1* and *KRT18* as well as downregulation of *VIM* (Fig. 5A). On the other hand, silencing *WT1* exacerbates their mesenchymal characteristics, with significant changes in the expression of EMT markers (Fig. 5A–C) along with a more elongated and irregular morphology, as shown by phalloidin staining (Fig. 6A). Moreover, even though no significant differences are observed at the mRNA level, the knockdown cells display nuclear accumulation of *CDH1* (Fig. 6C), a feature that has been linked to increased tumour invasion⁵⁰.

Some of these results, such as the upregulation of *TNC* upon *WT1* loss, were replicated in the other mesenchymal line used in this study, MDA-MB-231 (Figure S3 in Additional file 1). However, completely opposite results were observed in the epithelial line HBL100, where *WT1* overexpression led to the up-regulation of mesenchymal markers (Fig. 5B).

In order to understand which scenario more closely represents what happens *in vivo*, we compared the expression of EMT-related transcripts (*VIM*, *SNAI1*, *SNAI2*, *TWIST1*, *CDH1*, *CDH3*, *CLDN7*, *TNC*) between *WT1*-positive and *WT1*-negative samples from the integrated dataset described in Fig. 1A: statistically significant differences were only observed for the mesenchymal genes *VIM*, *SNAI1* and *TNC*, which showed higher expression in the *WT1*-positive samples (p -value < 0.001 , < 0.05 , < 0.001 respectively) (Fig. 5D, Figure S4 in Additional file 1).

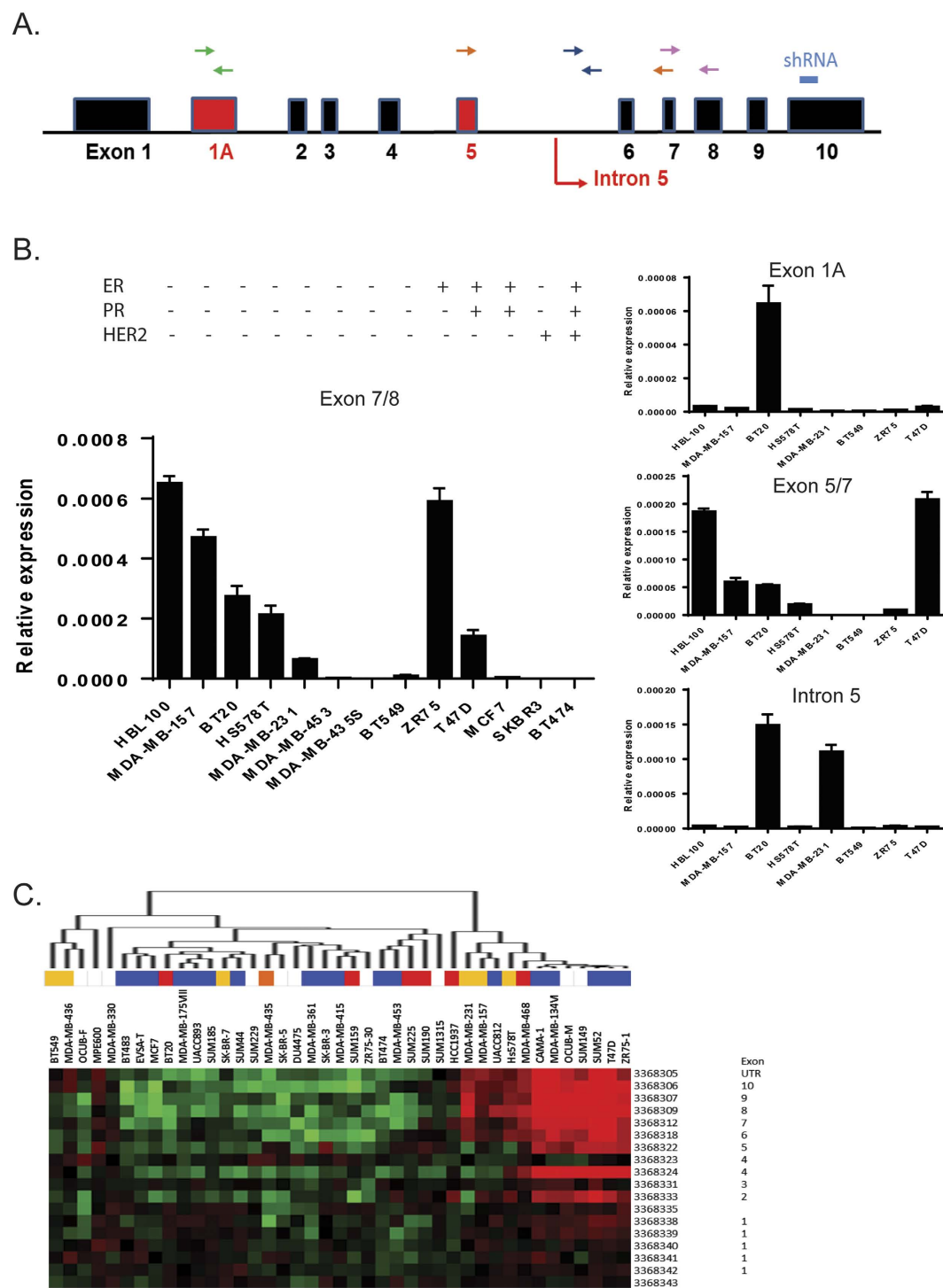


Figure 2. *WT1* expression in breast cancer cell lines. (A) Schematic of the *WT1* locus, showing alternative exons/splices in red, the position of the primers for each qRT-PCR assay and the sequence targeted by the shRNA used in the knockdown experiments. (B) Quantitative real-time PCR of *WT1* mRNA: data points represent the relative expression for each assay, error bars represent the standard deviation of two biological replicates. (C) Heatmap visualization of the relative expression of probes representing each *WT1* exon across a panel of breast cancer cell lines in a published dataset (GSE16732). The different colours indicate the cell line subtype (red = basal, orange = basal B/mesenchymal, purple = HER2 amplified, blue = luminal, yellow = claudin-low).

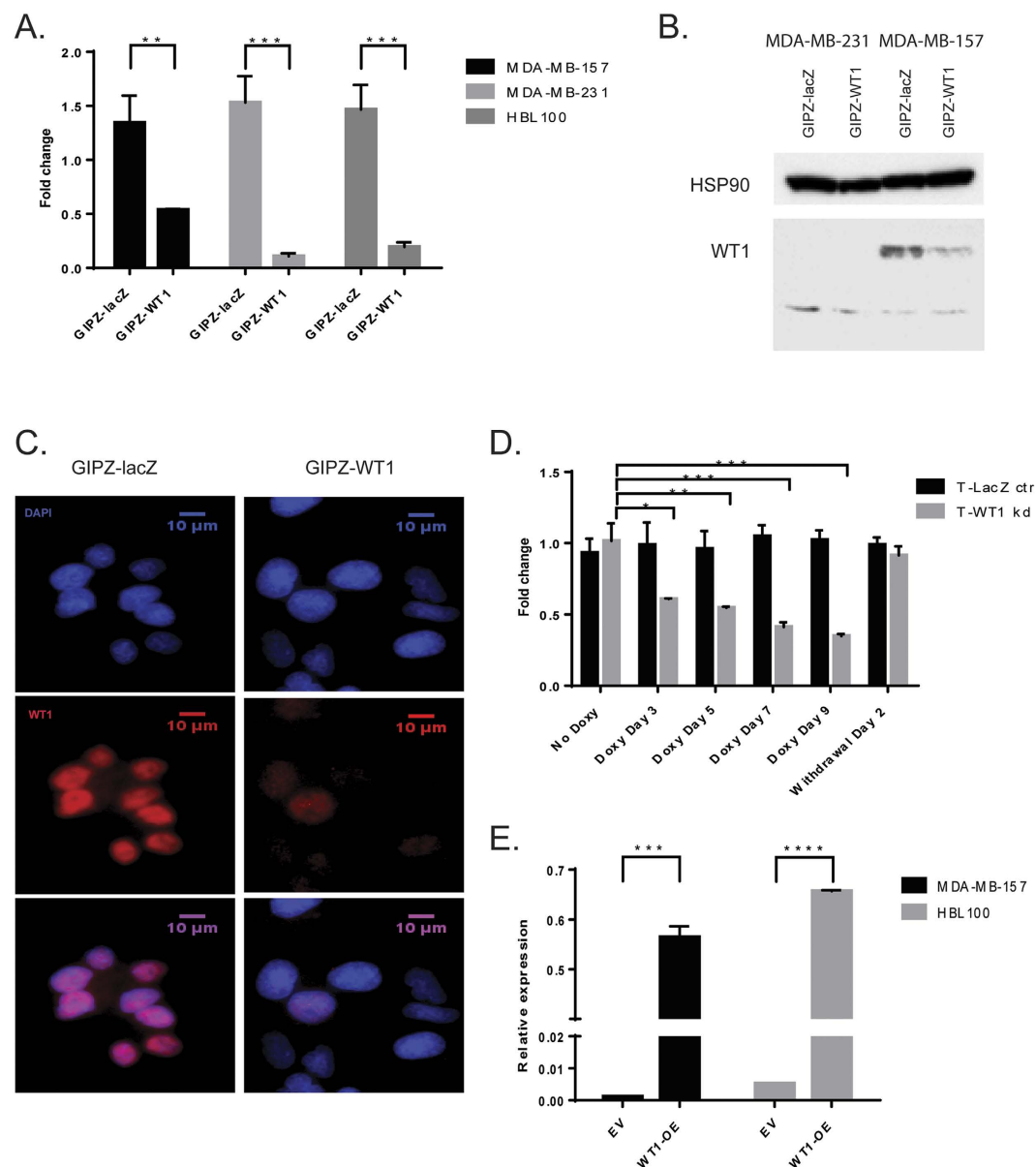


Figure 3. *WT1* knockdown and overexpression is achieved in breast cancer cell lines. (A) Quantitative real-time PCR of *WT1* mRNA in MDA-MB-157, MDA-MB-231 and HBL100 cells transduced with pGIPZ-miR: the graph represents fold change of mean expression relative to untransduced cells (given a value of 1); error bars represent the standard deviation of two separate biological replicates (** $p < 0.01$, *** $p < 0.001$). (B) Immunoblot of whole-protein lysates (40 μ g) probed with antibodies to *WT1* and HSP90, as a loading control. Only the truncated isoform of *WT1* can be observed in MDA-MB-231 cells. Full-length blots are presented in Figure S8 in Additional file 1. (C) *WT1* immunofluorescence of MDA-MB-157 cells transduced with pGIPZ-miR. (D) Quantitative real-time PCR of *WT1* mRNA in MDA-MB-157 cells transduced with pTRIPZ-miR: the cells were cultured on 2 μ g/ml doxycycline, the graph represents fold change of mean expression relative to untransduced cells, error bars represent the standard deviation of three biological replicates (* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$). (E) Quantitative real-time PCR of *WT1* mRNA in MDA-MB-157 and HBL100 cells transfected with pCI-WT1-OE; data points represent the relative expression, error bars represent the standard deviation of three biological replicates (** $p < 0.01$, **** $p < 0.0001$).

The results were further validated with the locally derived cohort previously described: the patient samples were first sorted according to their levels of *WT1* (Figure S1 in Additional file 1), then qPCR was used to assess the expression of the mesenchymal markers *VIM*, *SNAI1* and *TNC* (Fig. 5E). The *WT1*-positive tumours had higher expression of all three markers, even though statistical significance was only achieved for *TNC*.

Down-regulation of *WT1* does not result in changes in proliferation, apoptosis or cancer stem cell characteristics. Transient silencing of *WT1* has been previously shown to inhibit proliferation and induce apoptosis in cultures originated from leukemic cells and several solid tumours^{16–18,45}, therefore we also

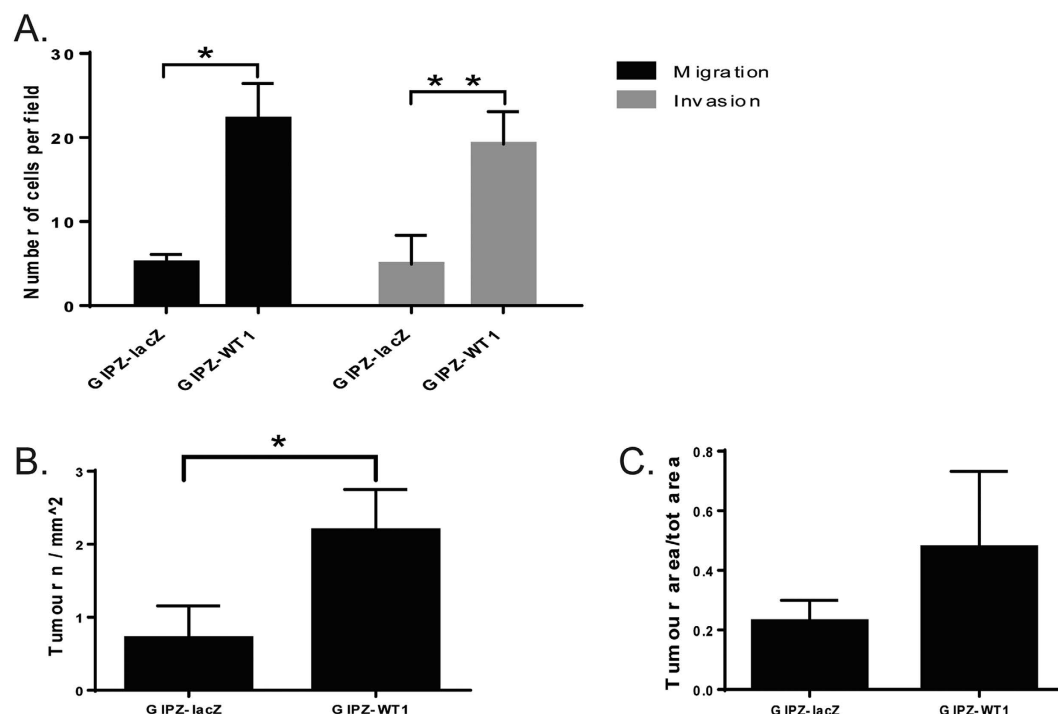


Figure 4. Knockdown of WT1 results in increased motility, invasiveness and metastatic potential of MDA-MB-157 cells. (A) Bar diagram representing the mean number of migrating and invading cells from three independent experiments. (B,C) WT1 knockdown significantly increases the number (B) but not the size (C) of lung metastases. MDA-MB-157 cells transduced with pGIPZ-miR were injected into nude mice through the tail vein and lung metastases were assessed 8 weeks after injection (n = 4 for each group).

performed apoptosis and cell cycle analysis. However, our knockdown cells did not show any significant change in either of these (Figures S5–6 in Additional file 1), suggesting that, at least for breast cancer cells with stable WT1 knockdown, the main phenotype is the disruption of the epithelial/mesenchymal balance, rather than apoptosis or cell cycle effects.

Since the EMT process has been linked to the acquisition of cancer stem-cell properties (e.g., an increased ability to form spheres in colony-forming assays or the expression of stem-cell markers)⁵¹, we also analysed stem-cell characteristics in the *WT1* knockdown cells.

No statistically significant difference in mammosphere formation was observed in any of the three lines used in this study (Fig. 7A). The cancer stem cell content of the knockdown cells was analysed by quantifying the sub-population of CD44+/CD24– cells, which has been reported to be highly enriched in CSCs⁵¹. In the MDA-MB-157 line the percentage of CD44+/CD24– cells did not change significantly upon WT1 loss (Fig. 7B), suggesting that down-regulation of WT1 does not affect the ‘stemness’ of breast cancer cells.

WT1 expression correlates with CYP3A4 levels and is associated with poorer response to taxane treatment. In order to identify differentially expressed genes between *WT1*-positive and *WT1*-negative breast tumours, significance analysis of microarray (SAM) was performed using the publicly available datasets previously described⁴¹. Among the 500 most significantly differentially expressed transcripts (Table S3 in Additional file 2) there are several different types of metalloproteases and many genes belonging to the cytochrome P450 family (CYP), all of which are significantly ($p < 0.0001$) positively correlated with WT1 (Fig. 8A).

These oxidising enzymes play an important and complex role in the metabolism of anti-cancer agents: they participate in the biological activation of many pro-drugs, but also to the inactivation of some cytotoxic compounds, two processes that occur predominantly, but not exclusively, in the liver. Not surprisingly, polymorphisms and different expression levels of the CYP genes have been shown to influence the outcome of chemotherapeutic treatments⁵².

The substrates of one of these up-regulated genes, *CYP3A4*, include drugs routinely used in the management of breast cancer such as tamoxifen, docetaxel, paclitaxel, cyclophosphamide and doxorubicin⁵². More specifically, *CYP3A4* activity in the liver is responsible for the activation of tamoxifen and cyclophosphamide, whereas it transforms docetaxel into inactive hydroxylated derivatives; interestingly, high *CYP3A4* levels in breast cancer biopsies can predict a poor therapeutic response to docetaxel, which suggests that also the enzyme within the tumour is involved in the inactivation of this drug⁵³.

Given this association between *WT1* and *CYP3A4* expression in breast cancer (Fig. 8B), we decided to investigate whether the tumour levels of *WT1* can be used to predict the response of breast cancer patients to docetaxel. To do so, we analysed a publicly available gene expression dataset (GSE25065) of 198 HER2-negative breast cancer cases treated with taxane-anthracycline chemotherapy pre-operatively and endocrine therapy if ER-positive⁵⁴.

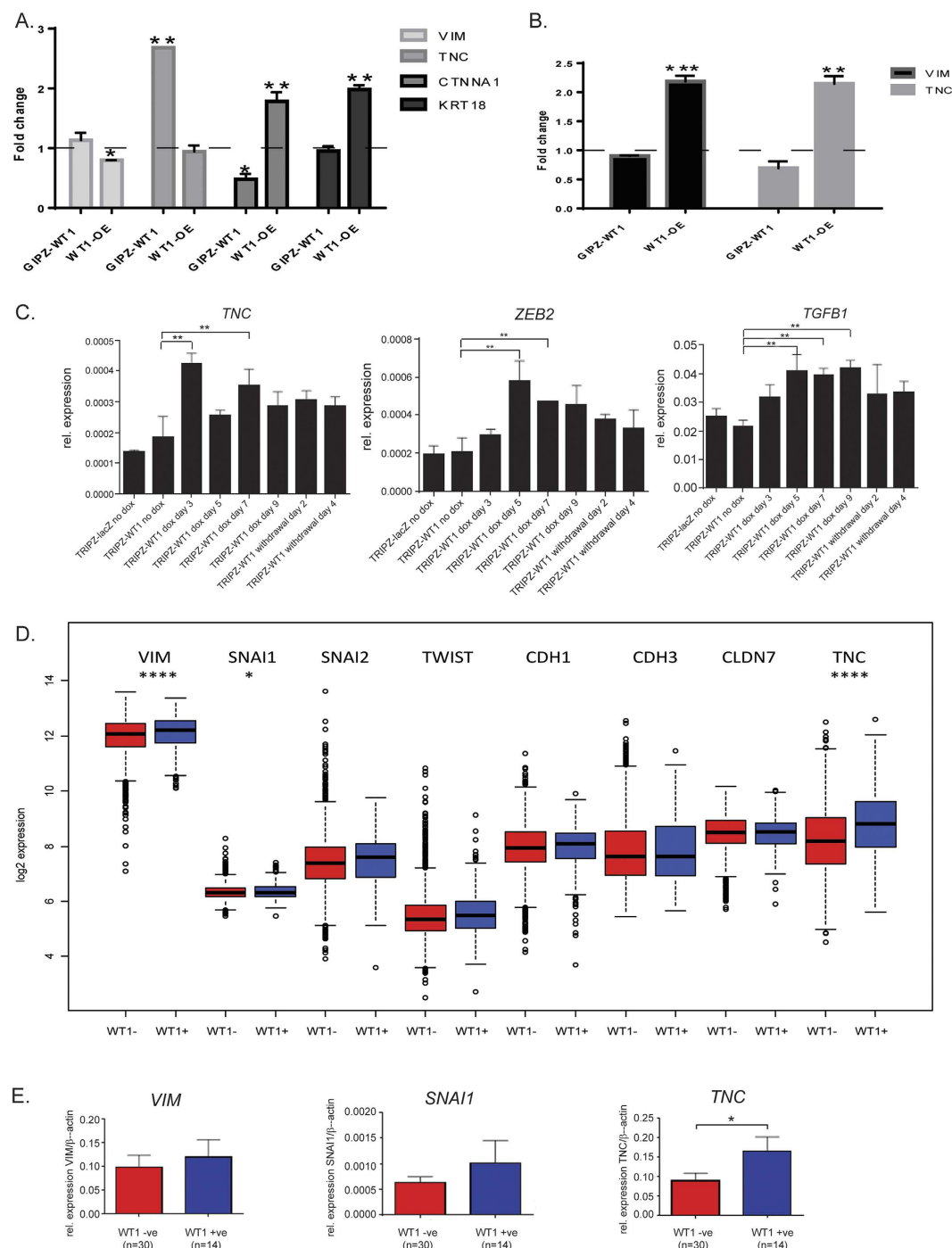


Figure 5. Correlation between WT1 expression and EMT markers *in vitro* and *in vivo*. (A) Quantitative real-time PCR of epithelial and mesenchymal markers in MDA-MB-157 cells transduced with pGIPZ-miR or transfected with pCI-WT1-OE: the graph represents fold change of mean expression relative to control cells, error bars represent the standard deviation of three biological replicates (* $p < 0.05$, ** $p < 0.01$). (B) Quantitative real-time PCR of mesenchymal markers in HBL100 cells transduced with pGIPZ-miR or transfected with pCI-WT1-OE: the graph represents fold change of mean expression relative to control cells, error bars represent the standard deviation of three biological replicates (** $p < 0.01$, *** $p < 0.001$). (C) Quantitative real-time PCR of *TNC*, *TGFβ1* and *ZEB2* mRNA in MDA-MB-157 cells transduced with pTRIPZ-miR: data points represent the relative expression of the gene, error bars represent the standard deviation of three separate biological replicates (** $p < 0.01$). (D) Boxplot of EMT genes expression in the WT1-positive and WT1-negative samples obtained from 17 integrated datasets ($n = 2999$ tumours), y-axis showing log2 values. Statistically significant differences were observed only for *VIM*, *SNAIL1* and *TNC*, which showed higher expression in the WT1-positive samples. * $p < 0.05$, *** $p < 0.0001$. (E) Quantitative RT-PCR of EMT genes in clinical samples ($n = 44$ tumours).

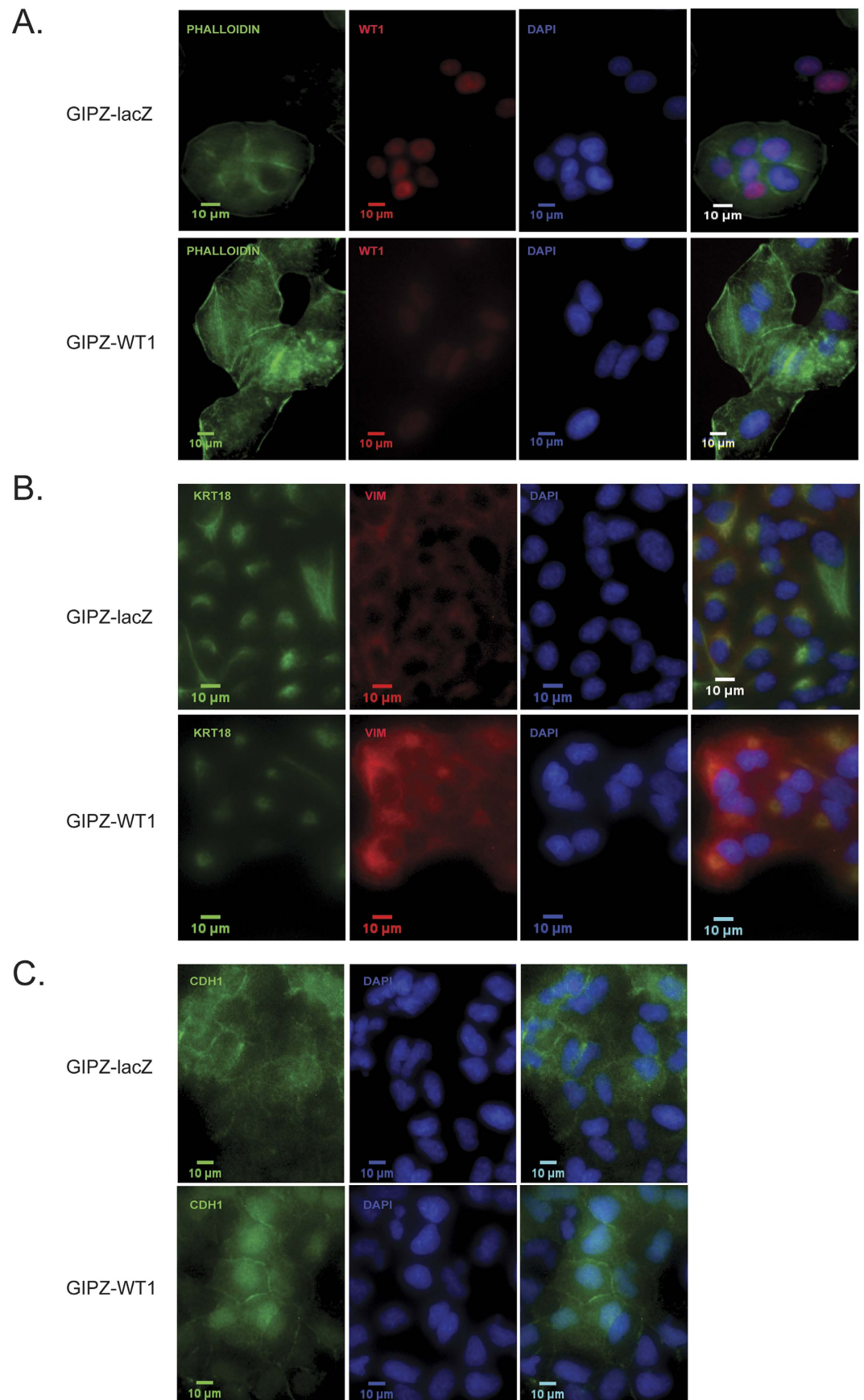


Figure 6. EMT markers immunofluorescence in MDA-MB-157 cells. MDA-MB-157 cells transduced with pGIPZ-miR were stained with phalloidin toxin (A), vimentin/cytokeratin18 (B) and E-cadherin (C) antibodies.

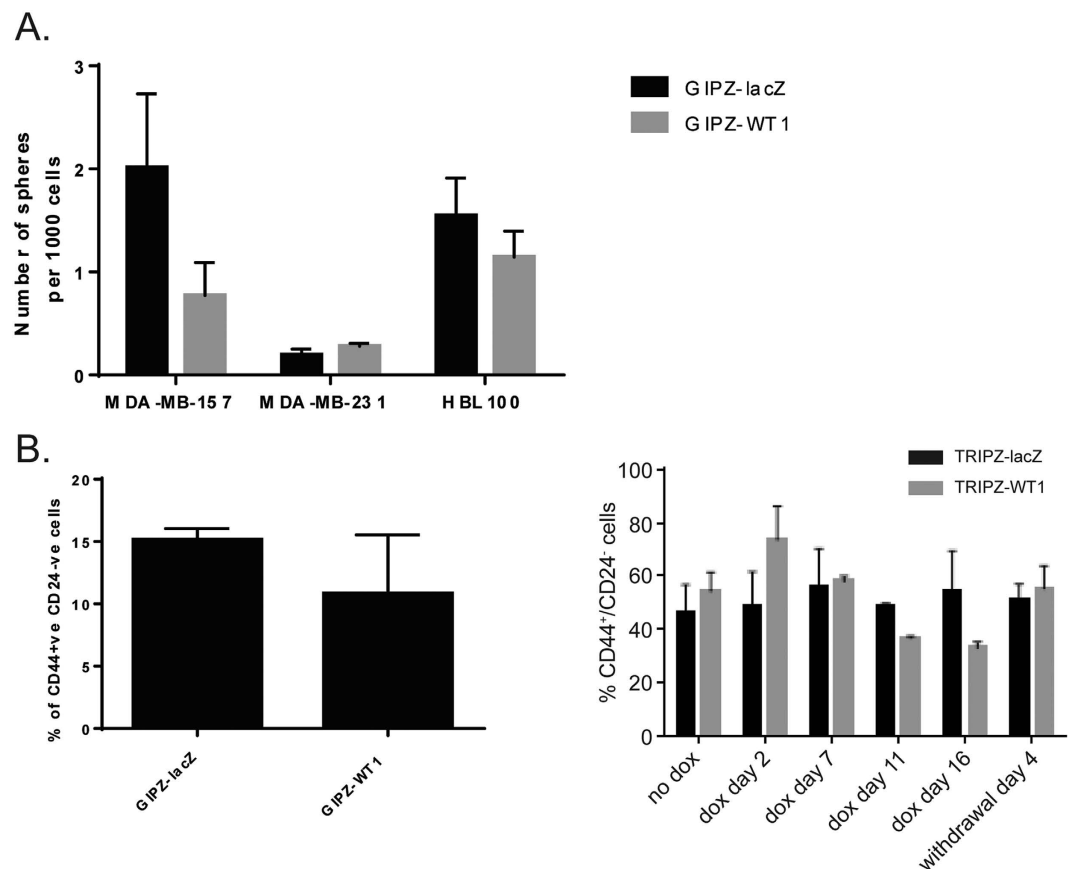


Figure 7. Knockdown of WT1 does not affect CSCs percentage or colony forming ability. (A) Colony forming assay of MDA-MB-157, MDA-MB-231 and HBL100 cells with constitutive WT1 knockdown suspended in semi-solid medium. Data points represent the average number of spheres formed per well, error bars represent the SEM of four biological replicates. (B) FACS analysis of CD24 and CD44 expression in MDA-MB-157 cells transduced with pGIPZ-miR and pTRIZ-miR. Bar graphs represent the average percentage of CD44+/CD24− cells in the different clones, error bars represent the SEM of three biological replicates.

As shown in Fig. 8C, *WT1* expression is significantly associated with poorer response and survival, with the 5-year recurrence-free percentage in the *WT1*-positive patients being 20% lower than in the *WT1*-negative (p value = 0.0072). To further validate these results, *CYP3A4* expression levels were analysed in the MDA-MB-157 line and showed significant down-regulation upon *WT1* loss (p = 0.035) (Fig. 8D); the *WT1* knockdown cells also displayed increased sensitivity to docetaxel if compared to GIPZ-lacZ control (Fig. 8E).

Discussion

Despite extensive studies, the role of *WT1* in breast cancer and its significance in terms of prognosis remain unclear, with conflicting results being reported in the literature.

Since this malignancy is a very heterogeneous disease and the *WT1* locus is extremely complex, we started our study with a comprehensive assessment of *WT1* expression in the different tumour subtypes followed by an isoform specific expression analysis.

Our work shows that *WT1* expression in breast cancer occurs at low frequency (10–30%), and it is lower than in the healthy mammary gland, which is in agreement with some of the earlier studies on the topic^{5,6}. Also, *WT1* expression is significantly higher in ER-positive (luminal) than in ER-negative (basal) tumours. This finding may be of particular interest, given that *WT1* has been shown to both interact with ER α and modulate its expression *in vitro*⁵⁵.

In terms of isoform expression pattern, all our data corroborate the hypothesis that the truncated transcript starting from intron5 is a tumour-specific isoform: this short *WT1* variant was first detected in the blood of patients affected by acute leukaemia, then in breast and pancreatic cancer cell lines^{43,56} and we have now shown that it is expressed in human breast cancer biopsies, but is absent in the healthy mammary gland.

Both our *in vitro* and *in silico* experiments also seem to demonstrate that the truncated intron5 is the only *WT1* isoform expressed in MDA-MB-231 cells; if confirmed, any assay targeted against exon1/5 would give a negative result, which may explain why the reports on *WT1* expression in this cell line are so contradictory^{7,24,45,57,58}.

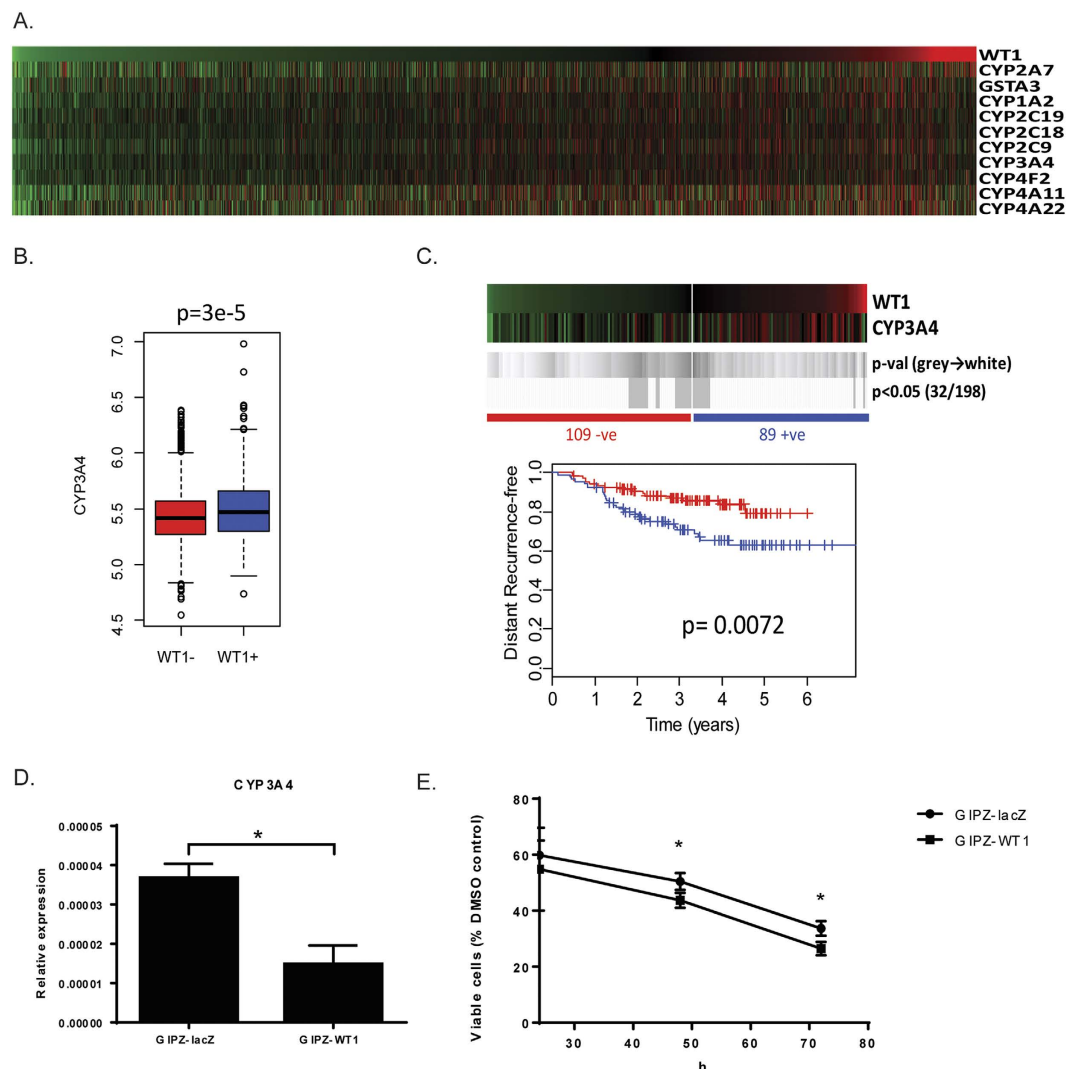


Figure 8. *WT1* expression in breast cancer correlates with cytochrome P450 family members and poor response to taxane treatment. (A) Heatmap showing ranked expression of *WT1* alongside cytochrome P450 genes obtained from 17 integrated published microarray datasets (n = 2999 tumours). (B) Boxplot of *CYP3A4* expression in the *WT1*-positive and *WT1*-negative samples. (C) Kaplan Meier survival analysis of 198 breast cancer patients treated with taxane-anthracycline chemotherapy (Dataset GSE25065) demonstrates that *WT1* expression is associated with worse outcomes. Heatmaps show log2 mean-centered expression values, red = high, green = low. Shades of grey to white indicate p-values of log-rank (Mantel-Cox) tests at all possible cut-points are shown in grey. Vertical bars on survival curves indicate censored cases. (D) Quantitative real-time PCR of *CYP3A4* mRNA in MDA-MB-157 cells transduced with pGIPZ-miR: data points represent the relative expression of the gene, error bars represent the standard deviation of two separate biological replicates ($*p < 0.05$). (E) Cytotoxicity of docetaxel is increased in MDA-MB-157 cells transduced with pGIPZ-miR. Mean and standard deviation of three independent experiments are plotted ($*p < 0.05$).

The last isoform we analysed, derived from a maternally imprinted alternative exon¹⁴², has been detected in few human breast cancer samples and in only one cell line, suggesting that it is unlikely to play a major role in breast tumorigenesis.

WT1 expression in invasive ductal carcinoma has been associated with both improved¹⁴ and worse⁸ outcomes, while a third study, specifically on triple negative breast cancers¹⁵ has shown that high *WT1* levels correlate with poor survival. *WT1* expression in other adult tumours is usually associated with higher histological grade as well as worse prognosis^{59,60} and several hypotheses have been formulated to explain this association, such as altered proliferation/apoptosis^{16–19}, increased angiogenesis^{25–27} and induction of cancer-EMT¹.

In this study, we focused our analysis on the relationship between *WT1* expression and the overall epithelial/mesenchymal balance of breast cancer cells by developing the first stable *WT1* knockdown in human breast cancer cells and extensively studying the effects of *WT1* silencing as well as overexpression.

When we perturb the expression levels of *WT1*, either by knocking it down or over-expressing it, the breast cancer cells show a disruption of their epithelial-mesenchymal balance, which seems to be dependent on their

baseline characteristics. In particular, in mesenchymal-leaning lines such as MDA-MB-157 and MDA-MB-231, the loss of WT1 pushes them towards an even less epithelial phenotype, with increased motility, invasiveness and metastatic potential (as shown by the tail vein injection experiment). The opposite holds true for the epithelial line HBL100, where it is WT1 over-expression that leads to a more mesenchymal state. Such a dichotomous role is not surprising for WT1, which is known to regulate both EMT and MET in development^{2,3}.

In terms of gene expression, the most significant changes are observed for the epithelial marker *CTNNA1* (a catenin that maintains normal cell adhesion by forming a complex that anchors E-cadherin to the cell membrane and if disrupted can lead to tumour invasion) and the mesenchymal marker *TNC* (an extracellular matrix glycoprotein which is usually found at the invasive front of the tumour⁶¹ and has been shown to induce EMT-like changes and faster migration in breast cancer cells⁶²). Interestingly, *VEGF*, which is an established WT1 target²⁸ is likely upstream of *TNC*⁶³.

Overall, perturbing the expression levels of *WT1* affects the epithelial/mesenchymal balance of cancer cells, but does not cause any significant change in cancer stem-cell properties, cell proliferation or apoptosis.

This finding is apparently in contrast with some previous experiments on leukemic and solid tumours cell lines, where the transient silencing of *WT1* led to growth arrest and apoptosis^{16–18,45}. However, the activation of an EMT program has been shown to protect from pro-apoptotic signals⁶⁴; this could explain the discrepancy for MDA-MB-157 and MDA-MB-231 cells, which increase their mesenchymal features upon WT1 loss.

The *in vitro* experiments were then corroborated by the analysis of publicly available gene expression datasets of breast cancer cell lines and clinical samples: overall, the results are consistent with what observed in the HBL100 cell line and suggest that *WT1* expression is usually associated with a more mesenchymal phenotype.

The gene expression analysis also revealed that the *WT1*-positive tumours exhibit poorer response to taxane treatment, which may be linked to their increased levels of *CYP3A4*, an oxidising enzyme which transforms docetaxel into inactive hydroxylated derivatives⁶⁵. Importantly, *CYP3A4*, whose expression levels correlate with *WT1* both *in vivo* and *in vitro*, is activated by VDR⁶⁶, a known WT1 target⁶⁷.

As a whole, this study has shown that WT1 can disrupt the epithelial-mesenchymal balance of breast cancer lines in opposite ways, depending on the baseline characteristics of the cells.

In breast tumours however, the *WT1*-expressing tumours are mainly associated with a mesenchymal phenotype and high levels of *CYP3A4*: therefore, their cells not only respond poorly to taxane, but they are also more likely to invade the blood vessels and metastasise.

In conclusion, we demonstrate that cancer-related EMT and poor chemotherapy response are likely the main factors that explain why WT1 expression in breast cancer is usually associated with poor prognosis.

Methods

Cell culture. The human breast cancer cell lines MDA-MB-157, MDA-MB-231, MDA-MB-453, MDA-MB-435S, HBL100, MCF7, BT20, SKBR3, T47D, HS578T and ZR75 were maintained in DMEM medium (Sigma); BT474 and BT549 cell lines in RPMI medium (Sigma). Both media were supplemented with 10% fetal bovine serum (FBS), 2 mM L-glutamine, 100 units/mL penicillin and 100 mg/mL streptomycin. The human embryonic kidney cell line SODK3 used for the lentiviral packaging was maintained in DMEM-HIGH (Sigma) supplemented with 10% FBS, 2 mM L-glutamine, 100 units/mL penicillin, 100 mg/mL streptomycin, 1 µg/mL puromycin, 600 µg/mL neomycin and 1 µg/mL doxycycline. All lines were kept at standard cell culture conditions (37°C, 5% CO₂ in humidified incubator). Tet System Approved FBS (Clontech) was used in the culture media of TRIPZ-transduced cells, to ensure that the inducible regulation of the system was not altered by the presence tetracycline-derived contaminants.

Clinical samples. The use of tissue from invasive breast cancer patients treated at the Edinburgh Breast Unit at the Western General Hospital was approved by the Lothian Research Ethics Committee (08/S1101/41). This study was carried out in accordance with the approved guidelines and written informed consent was obtained from all subjects. Multiple core biopsies were harvested from consenting patients at the time of curative surgical resection for invasive breast cancer. Macroscopically distinct fat was trimmed and discarded. Tumour samples were then stored at −80°C until use.

shRNA-based stable WT1 knockdown and transient WT1 overexpression. The lentiviral GIPZ-shRNA^{mir}-GFP and TRIPZ-shRNA^{mir}-RFP vectors (Thermo Scientific Open Biosystems) were used as backbones for the cloning of shRNAs targeting the human *WT1* transcript and a lacZ control (Figure S7 in Additional file 1, Table S4 in Additional file 2).

The final constructs were transduced using a trans-lentiviral packaging system developed in the human embryonic kidney cell line SODK3⁶⁸, which expresses GFP and all trans-lentivirus packaging proteins from a Tet-Off system. To activate the packaging system, the cells were grown for 3 days without doxycycline and FACS sorted for GFP expression; the brightest 15% was plated on a 0.001% poly-L-Lysine coated plate and grown overnight without doxycycline. The following day, the SODK3 cells were transfected with the lentiviral vectors using FuGene6 (Roche) as per manufacturer's instructions; the medium, containing 5 mM sodium butyrate (Sigma) but no doxycycline, was changed at 12 and 36 hours post-transfection. 72 hours after the last medium change, the virus-containing supernatants were harvested and centrifuged for 20 minutes at 4°C to pellet cell debris. Standard procedures were used to transduce MDA-MB-157, MDA-MB-231 and HBL100 cells, which were maintained on selective media (1 µg/mL puromycin) from day 2 post-transduction.

Full-length WT1 cDNA was amplified and cloned into the pCI mammalian expression vector using In-Fusion (Clontech). Cells were transfected with pCI-WT1-OE or the empty vector (Figure S7 in Additional file 1) and harvested after 30 h.

Cytotoxicity assay. Cells were plated onto 96-well plates (5×10^3 cells/well) and after 24 hours the culture medium was replaced by fresh medium containing 40 nM docetaxel or DMSO. The cells were incubated for another 24, 48 and 72 h, then washed with PBS, fixed in cold methanol and stained with 0.5% crystal violet for 30 minutes. After adding 100 μ l Sorenson buffer/well, absorbance was measured as a read-out of viable cells.

Animal experiments. Lung colonization experiments were performed in compliance with the UK Animals (Scientific Procedures) Act 1986, under Project Licence PPL 60/3788 approved by the UK Home Office. 1×10^6 cells suspended in 100 μ l sterile PBS were directly injected into the lateral tail vein of 4-week-old female athymic nude mice. Lung metastases were identified by histological staining of paraffin sections with hematoxylin and eosin. The slides were scanned with a Hamamatsu nanosizer slide scanner (Hamamatsu, UK), and then examined by two boarded pathologists -blinded to the experimental groups- for the presence of metastases using Hamamatsu NDP2 view software (Hamamatsu, UK). One representative section was chosen for each pulmonary lobe available for examination and measured. The measurements included total number of metastatic deposits, and the relative area of metastasis, expressed as total metastatic area/total lobe area for each sample.

Colony forming assay. Sphere formation was induced by culturing the cancer cells in suspension in serum-free medium (DMEM-F12 + GlutaMAX-I, Life Technologies) supplemented with B27 (1:50, Life Technologies), 20 ng/mL EGF (R&D), 0.4% bovine serum albumin (Sigma), 4 μ g/mL insulin (Sigma). To quantify colony formation, cells were plated at a density of 1,000 cells per well in 96-well plates coated with poly(2-hydroxyethyl metacrylate) to prevent cell attachment to the surface (Sigma); cells were diluted in supplemented serum-free media containing 1% methylcellulose (Sigma) and the number of colonies $>75 \mu$ m in diameter was counted after 11 days³⁸.

Immunofluorescence microscopy. Cells were fixed for 15 minutes in 4% paraformaldehyde, rinsed with PBS and permeabilised for 10 minutes in 0.1% Triton X-100. Nonspecific immunoglobulin binding was blocked with a 30 minute incubation in 3% bovine serum albumin + 0.1% Triton X-100. Primary antibodies (E-cadherin Cell Signalling 24E10, Vimentin Cell Signalling 5741P, Wt1 Abcam 89901, Cytokeratin18 Santa Cruz 31700) were diluted 1:100 in 1% bovine serum albumin + 0.1% Triton X-100. After overnight incubation at 4°C, cells were rinsed with PBS, then incubated for 1 h at room temperature with secondary antibodies (1:500 Alexa Fluor Invitrogen) or Phalloidin (Invitrogen A12379). Vectashield with DAPI (Vector Laboratories) was used to counterstain the nuclei and mount the slides.

Migration and invasion assays. Migration assays were performed on 24 well plates containing 8 μ m PET culture inserts (Fisher Scientific). For each experiment, 5×10^4 cells resuspended in serum-free media were applied to the upper chamber while the lower chamber was filled with 300 μ l medium with 10% FBS. Cells were then incubated under normal conditions for 3.5 hours.

The invasion assay was similar to the migration assay, except that the inserts were evenly coated with 1 mg/ml Matrigel; 2.5×10^4 cells were added to the upper chamber and incubated for 24 h. For both assays, cells were fixed with methanol for 10 min and stained with 0.5% crystal violet for 30 min. After removing the cells on the upper side of the filters with cotton-tipped swabs, the cells on the underside of the filters were counted under a X20 objective lens in five randomly chosen fields.

Cancer stem cell markers, apoptosis and cell cycle analysis. For the analysis of cancer stem cell markers, 1×10^6 cells were resuspended in 100 μ l PBS/5% FBS and incubated in the antibody dilution shown in Table S5 in Additional File 2. The GIPZ-transduced cells were incubated with CD44-APC and CD24-PE antibodies, the TRIPZ-transduced cells with CD44-APC and CD24-FITC. After a 15 minute incubation on ice, the cells were washed twice in PBS, resuspended in 100 μ l PBS/5% FBS and analysed. The Annexin V Apoptosis Detection Kit APC (eBioscience) was used according to the manufacturer's instructions: cells were washed once and resuspended at a concentration of 1×10^6 cells/ml in the binding buffer provided with the kit; 5 μ l of the APC-conjugated antibody were added to 100 μ l of cell suspension and incubated for 15 minutes at room temperature. After one wash, cells were resuspended in 200 μ l of binding buffer and 1 μ g/ml DAPI (Sigma) was added immediately before the analysis. For cell cycle analysis, cells were trypsinised and resuspended in 1.2 ml PBS; 3 ml ice cold 95% ethanol were then added dropwise while vortexing, to obtain a final 70% ethanol solution where the cells were incubated for 30 minutes.

All experiments were performed on a FACSariaII Flow Cytometer (Becton Dickinson), data were analysed using FlowJo software.

RNA extraction, quantitative real-time PCR and RNA sequencing. Total RNA was extracted using RNeasy Mini Kit (Qiagen) with on-column DNase digestion as per manufacturer's instructions; cDNA was synthesized from 2 μ g total RNA using AMV reverse transcriptase (Roche) and oligo d(T) priming. All the quantitative Real Time PCR experiments were performed in triplicate on the Roche Lightcycler 480 system using Universal Probe Library probes and primers designed with the Universal Probe Library Assay Design Centre (Table S6 in Additional file 1). Relative quantification of gene expression was performed by calculating Δ Ct where Δ Ct = CtX - CtACTB. The fold change in gene expression between two samples was then determined by calculating $2^{-\Delta\Delta$ Ct. For RNA sequencing, total RNA was isolated as previously described and sent to GATC Biotech (Germany) for library preparation and sequencing on the Illumina HiSeq 2000 platform. The raw data were analysed using Galaxy (<https://usegalaxy.org/>) and have been deposited in NCBI's Gene Expression Omnibus⁶⁹, where they are accessible through GEO Series accession number GSE93636.

An RNA sample of human adult mammary gland was purchased from AMSBIO and retrotranscribed; the specimen consisted of a pool from 5 healthy donors of different age (36, 57, 76, 78 and 83 years old).

Gene expression analysis. Raw data (.cel) files from seventeen Affymetrix U133A/plus2 primary breast tumour gene expression datasets were downloaded from NCBI GEO (GSE12276, GSE21653, GSE3744, GSE5460, GSE2109, GSE1561, GSE17907, GSE2990, GSE7390, GSE11121, GSE16716, GSE2034, GSE1456, GSE6532, GSE3494) or the caBIG (geral-00143) repositories, normalised, batch-corrected and analysed as previously described⁴¹. The molecular subtypes were assigned based upon the highest correlation to the Prat *et al.*⁷⁰ centroids applied to each dataset separately. The Affymetrix U133A primary breast tumour gene expression dataset GSE25065⁵⁴ was used to compare the response to taxane-anthracycline chemotherapy in WT1-positive and WT1-negative tumours.

Immunoblot analysis. Immunoblotting was performed according to standard procedures using antibodies against WT1 (Novus Biologicals NBP1-40787, diluted 1/2000) and HSP90 (BD Bioscience 610419, diluted 1/5000).

References

- Hohenstein, P. & Hastie, N. D. The many facets of the Wilms' tumour gene, WT1. *Human molecular genetics* 15 Spec No 2, R196–201, doi: 10.1093/hmg/ddl196 (2006).
- Martinez-Estrada, O. M. *et al.* Wt1 is required for cardiovascular progenitor cell formation through transcriptional control of Snail and E-cadherin. *Nature genetics* 42, 89–93, doi: 10.1038/ng.494 (2010).
- Essafi, A. *et al.* A wt1-controlled chromatin switching mechanism underpins tissue-specific wnt4 activation and repression. *Developmental cell* 21, 559–574, doi: 10.1016/j.devcel.2011.07.014 (2011).
- Chau, Y. Y. & Hastie, N. D. The role of Wt1 in regulating mesenchyme in cancer, development, and tissue homeostasis. *Trends in genetics: TIG* 28, 515–524, doi: 10.1016/j.tig.2012.04.004 (2012).
- Silberstein, G. B., Van Horn, K., Strickland, P., Roberts, C. T. & Daniel, C. W. Altered expression of the WT1 Wilms tumor suppressor gene in human breast cancer. *Proceedings of the National Academy of Sciences of the United States of America* 94, 8132–8137 (1997).
- Cheng, C. W. *et al.* Mechanisms of inactivation of E-cadherin in breast carcinoma: modification of the two-hit hypothesis of tumor suppressor gene. *Oncogene* 20, 3814–3823, doi: 10.1038/sj.onc.1204505 (2001).
- Loeb, D. M. *et al.* Wilms' tumor suppressor gene (WT1) is expressed in primary breast tumors despite tumor-specific promoter methylation. *Cancer research* 61, 921–925 (2001).
- Miyoshi, Y. *et al.* High expression of Wilms' tumor suppressor gene predicts poor prognosis in breast cancer patients. *Clinical cancer research: an official journal of the American Association for Cancer Research* 8, 1167–1171 (2002).
- Silberstein, G. B., Dressler, G. R. & Van Horn, K. Expression of the PAX2 oncogene in human breast cancer and its role in progesterone-dependent mammary growth. *Oncogene* 21, 1009–1016, doi: 10.1038/sj.onc.1205172 (2002).
- Gillmore, R. *et al.* Detection of Wilms' tumor antigen-specific CTL in tumor-draining lymph nodes of patients with early breast cancer. *Clinical cancer research: an official journal of the American Association for Cancer Research* 12, 34–42, doi: 10.1158/1078-0432.ccr-05-1483 (2006).
- Nakatsuka, S. *et al.* Immunohistochemical detection of WT1 protein in a variety of cancer cells. *Modern pathology: an official journal of the United States and Canadian Academy of Pathology, Inc* 19, 804–814, doi: 10.1038/modpathol.3800588 (2006).
- Wang, L. & Wang, Z. Y. The Wilms' tumor suppressor WT1 inhibits malignant progression of neoplastigenic mammary epithelial cells. *Anticancer research* 28, 2155–2160 (2008).
- Burwell, E. A., McCarty, G. P., Simpson, L. A., Thompson, K. A. & Loeb, D. M. Isoforms of Wilms' tumor suppressor gene (WT1) have distinct effects on mammary epithelial cells. *Oncogene* 26, 3423–3430, doi: http://www.nature.com/onc/journal/v26/n23/supinfo/1210127s1.html (2006).
- Choi, E. J. *et al.* Prognostic significance of RSPO1, WNT1, P16, WT1, and SDC1 expressions in invasive ductal carcinoma of the breast. *World J Surg Oncol* 11, 314, doi: 10.1186/1477-7819-11-314 (2013).
- Stirzaker, C. *et al.* Methylation sequencing in triple-negative breast cancer reveals distinct methylation clusters with prognostic value. *Nat Commun* 6, 5899, doi: 10.1038/ncomms6899 (2015).
- Yamagami, T. *et al.* Growth inhibition of human leukemic cells by WT1 (Wilms tumor gene) antisense oligodeoxynucleotides: implications for the involvement of WT1 in leukemogenesis. *Blood* 87, 2878–2884 (1996).
- Algar, E. M. *et al.* A WT1 antisense oligonucleotide inhibits proliferation and induces apoptosis in myeloid leukaemia cell lines. *Oncogene* 12, 1005–1014 (1996).
- Zamora-Avila, D. E. *et al.* RNAi silencing of the WT1 gene inhibits cell proliferation and induces apoptosis in the B16F10 murine melanoma cell line. *Melanoma research* 17, 341–348, doi: 10.1097/CMR.0b013e3282ef3ae (2007).
- Tatsumi, N. *et al.* Wilms' tumor gene WT1-shRNA as a potent apoptosis-inducing agent for solid tumors. *International journal of oncology* 32, 701–711 (2008).
- Hewitt, S. M., Hamada, S., McDonnell, T. J., Rauscher, F. J., 3rd & Saunders, G. F. Regulation of the proto-oncogenes bcl-2 and c-myc by the Wilms' tumor suppressor gene WT1. *Cancer research* 55, 5386–5389 (1995).
- Morrison, D. J., English, M. A. & Licht, J. D. WT1 induces apoptosis through transcriptional regulation of the proapoptotic Bcl-2 family member Bak. *Cancer research* 65, 8174–8182, doi: 10.1158/0008-5472.can-04-3657 (2005).
- Cesaro, E. *et al.* WT1 protein is a transcriptional activator of the antiapoptotic bag3 gene. *Leukemia* 24, 1204–1206, doi: 10.1038/leu.2010.68 (2010).
- Uesugi, K. *et al.* Wilms' tumor 1 gene modulates Fas-related death signals and anti-apoptotic functions in hepatocellular carcinoma. *Journal of gastroenterology* 48, 1069–1080, doi: 10.1007/s00535-012-0708-7 (2013).
- Caldon, C. E., Lee, C. S., Sutherland, R. L. & Musgrove, E. A. Wilms' tumor protein 1: an early target of progesterone regulation in T-47D breast cancer cells that modulates proliferation and differentiation. *Oncogene* 27, 126–138, doi: 10.1038/sj.onc.1210622 (2008).
- Timar, J., Meszaros, L., Orosz, Z., Albini, A. & Raso, E. WT1 expression in angiogenic tumours of the skin. *Histopathology* 47, 67–73, doi: 10.1111/j.1365-2559.2005.02169.x (2005).
- Wagner, N., Michiels, J. F., Schedl, A. & Wagner, K. D. The Wilms' tumour suppressor WT1 is involved in endothelial cell proliferation and migration: expression in tumour vessels *in vivo*. *Oncogene* 27, 3662–3672, doi: 10.1038/sj.onc.1211044 (2008).
- Dohi, S. *et al.* WT1 expression correlates with angiogenesis in endometrial cancer tissue. *Anticancer research* 30, 3187–3192 (2010).
- Hanson, J., Gorman, J., Reese, J. & Fraizer, G. Regulation of vascular endothelial growth factor, VEGF, gene promoter by the tumor suppressor, WT1. *Frontiers in bioscience: a journal and virtual library* 12, 2279–2290 (2007).
- Amin, E. M. *et al.* WT1 mutants reveal SRPK1 to be a downstream angiogenesis target by altering VEGF splicing. *Cancer cell* 20, 768–780, doi: 10.1016/j.ccr.2011.10.016 (2011).
- Kirschner, K. M., Sciesielski, L. K. & Scholz, H. Wilms' tumour protein Wt1 stimulates transcription of the gene encoding vascular endothelial cadherin. *Pflügers Archiv: European journal of physiology* 460, 1051–1061, doi: 10.1007/s00424-010-0873-6 (2010).
- Wagner, K. D. *et al.* The Wilms' tumour suppressor Wt1 is a major regulator of tumour angiogenesis and progression. *Nat Commun* 5, 5852, doi: 10.1038/ncomms6852 (2014).
- Fischer, K. R. *et al.* Epithelial-to-mesenchymal transition is not required for lung metastasis but contributes to chemoresistance. *Nature* 527, 472–476, doi: 10.1038/nature15748 (2015).

33. Zheng, X. *et al.* Epithelial-to-mesenchymal transition is dispensable for metastasis but induces chemoresistance in pancreatic cancer. *Nature* **527**, 525–530, doi: 10.1038/nature16064 (2015).
34. Trimboli, A. J. *et al.* Direct evidence for epithelial-mesenchymal transitions in breast cancer. *Cancer research* **68**, 937–945, doi: 10.1158/0008-5472.can-07-2148 (2008).
35. Brabletz, T., Jung, A., Spaderna, S., Hlubek, F. & Kirchner, T. Opinion: migrating cancer stem cells - an integrated concept of malignant tumour progression. *Nature reviews. Cancer* **5**, 744–749, doi: 10.1038/nrc1694 (2005).
36. Blanco, M. J. *et al.* Correlation of Snail expression with histological grade and lymph node status in breast carcinomas. *Oncogene* **21**, 3241–3246, doi: 10.1038/sj.onc.1205416 (2002).
37. Yang, J. *et al.* Twist, a master regulator of morphogenesis, plays an essential role in tumor metastasis. *Cell* **117**, 927–939, doi: 10.1016/j.cell.2004.06.006 (2004).
38. Wellner, U. *et al.* The EMT-activator ZEB1 promotes tumorigenicity by repressing stemness-inhibiting microRNAs. *Nature cell biology* **11**, 1487–1495, doi: 10.1038/ncb1998 (2009).
39. Dey, B. R. *et al.* Repression of the transforming growth factor-beta 1 gene by the Wilms' tumor suppressor WT1 gene product. *Molecular endocrinology (Baltimore, Md.)* **8**, 595–602, doi: 10.1210/mend.8.5.8058069 (1994).
40. Takeichi, M., Nimura, K., Mori, M., Nakagami, H. & Kaneda, Y. The transcription factors Tbx18 and Wt1 control the epicardial epithelial-mesenchymal transition through bi-directional regulation of Slug in murine primary epicardial cells. *PLoS one* **8**, e57829, doi: 10.1371/journal.pone.0057829 (2013).
41. Moleirinho, S. *et al.* KIBRA exhibits MST-independent functional regulation of the Hippo signaling pathway in mammals. *Oncogene* **32**, 1821–1830, doi: 10.1038/onc.2012.196 (2013).
42. Dallosso, A. R. *et al.* Genomic imprinting at the WT1 gene involves a novel coding transcript (AWT1) that shows deregulation in Wilms' tumours. *Human molecular genetics* **13**, 405–415, doi: 10.1093/hmg/ddh038 (2004).
43. Dechsukhum, C., Ware, J. L., Ferreira-Gonzalez, A., Wilkinson, D. S. & Garrett, C. T. Detection of a novel truncated WT1 transcript in human neoplasia. *Molecular diagnosis: a journal devoted to the understanding of human disease through the clinical application of molecular biology* **5**, 117–128, doi: 10.1054/xd.2000.7181 (2000).
44. Riaz, M. *et al.* Low-risk susceptibility alleles in 40 human breast cancer cell lines. *BMC cancer* **9**, 236, doi: 10.1186/1471-2407-9-236 (2009).
45. Zapata-Benavides, P., Tuna, M., Lopez-Berestein, G. & Tari, A. M. Downregulation of Wilms' tumor 1 protein inhibits breast cancer proliferation. *Biochemical and biophysical research communications* **295**, 784–790 (2002).
46. Li, X., Law, J. W. & Lee, A. Y. Semaphorin 5A and plexin-B3 regulate human glioma cell motility and morphology through Rac1 and the actin cytoskeleton. *Oncogene* **31**, 595–610, doi: 10.1038/onc.2011.256 (2012).
47. Gutschner, T. *et al.* The noncoding RNA MALAT1 is a critical regulator of the metastasis phenotype of lung cancer cells. *Cancer research* **73**, 1180–1189, doi: 10.1158/0008-5472.can-12-2850 (2013).
48. Chen, J., Bardes, E. E., Aronow, B. J. & Jegga, A. G. ToppGene Suite for gene list enrichment analysis and candidate gene prioritization. *Nucleic acids research* **37**, W305–311, doi: 10.1093/nar/gkp427 (2009).
49. Xie, G. *et al.* Tumour-initiating capacity is independent of epithelial-mesenchymal transition status in breast cancer cell lines. *British journal of cancer* **110**, 2514–2523, doi: 10.1038/bjc.2014.153 (2014).
50. Elston, M. S. *et al.* Nuclear accumulation of e-cadherin correlates with loss of cytoplasmic membrane staining and invasion in pituitary adenomas. *The Journal of clinical endocrinology and metabolism* **94**, 1436–1442, doi: 10.1210/jc.2008-2075 (2009).
51. Mani, S. A. *et al.* The epithelial-mesenchymal transition generates cells with properties of stem cells. *Cell* **133**, 704–715, doi: 10.1016/j.cell.2008.03.027 (2008).
52. Rodriguez-Antona, C. & Ingelman-Sundberg, M. Cytochrome P450 pharmacogenetics and cancer. *Oncogene* **25**, 1679–1691, doi: 10.1038/sj.onc.1209377 (2006).
53. Miyoshi, Y. *et al.* Prediction of response to docetaxel by CYP3A4 mRNA expression in breast cancer tissues. *International journal of cancer. Journal international du cancer* **97**, 129–132 (2002).
54. Hatzis, C. *et al.* A genomic predictor of response and survival following taxane-anthracycline chemotherapy for invasive breast cancer. *JAMA* **305**, 1873–1881, doi: 10.1001/jama.2011.593 (2011).
55. Han, Y. *et al.* Wilms' tumor 1 suppressor gene mediates antiestrogen resistance via down-regulation of estrogen receptor- α expression in breast cancer cells. *Molecular cancer research: MCR* **6**, 1347–1355, doi: 10.1158/1541-7786.mcr-07-2179 (2008).
56. Bae, V. L., Jackson-Cook, C. K., Brothman, A. R., Maygarden, S. J. & Ware, J. L. Tumorigenicity of SV40 T antigen immortalized human prostate epithelial cells: association with decreased epidermal growth factor receptor (EGFR) expression. *International journal of cancer. Journal international du cancer* **58**, 721–729 (1994).
57. Laux, D. E., Curran, E. M., Welshons, W. V., Lubahn, D. B. & Huang, T. H. Hypermethylation of the Wilms' tumor suppressor gene CpG island in human breast carcinomas. *Breast cancer research and treatment* **56**, 35–43 (1999).
58. Tuna, M., Chavez-Reyes, A. & Tari, A. M. HER2/neu increases the expression of Wilms' Tumor 1 (WT1) protein to stimulate S-phase proliferation and inhibit apoptosis in breast cancer cells. *Oncogene* **24**, 1648–1652, doi: 10.1038/sj.onc.1208345 (2005).
59. Oji, Y. *et al.* Overexpression of the Wilms' tumor gene WT1 in head and neck squamous cell carcinoma. *Cancer science* **94**, 523–529 (2003).
60. Ohno, S. *et al.* Immunohistochemical detection of WT1 protein in endometrial cancer. *Anticancer research* **29**, 1691–1695 (2009).
61. Jahkola, T. *et al.* Tenascin-C expression in invasion border of early breast cancer: a predictor of local and distant recurrence. *British journal of cancer* **78**, 1507–1513 (1998).
62. Nagaharu, K. *et al.* Tenascin C induces epithelial-mesenchymal transition-like change accompanied by SRC activation and focal adhesion kinase phosphorylation in human breast cancer cells. *The American journal of pathology* **178**, 754–763, doi: 10.1016/j.ajpath.2010.10.015 (2011).
63. Calvo, A. *et al.* Identification of VEGF-regulated genes associated with increased lung metastatic potential: functional involvement of tenascin-C in tumor growth and lung metastasis. *Oncogene* **27**, 5373–5384, doi: 10.1038/onc.2008.155 (2008).
64. Vega, S. *et al.* Snail blocks the cell cycle and confers resistance to cell death. *Genes & development* **18**, 1131–1143, doi: 10.1101/gad.294104 (2004).
65. Engels, F. K. *et al.* Effect of cytochrome P450 3A4 inhibition on the pharmacokinetics of docetaxel. *Clinical pharmacology and therapeutics* **75**, 448–454, doi: 10.1016/j.clpt.2004.01.001 (2004).
66. Pavik, P. *et al.* Intestinal cell-specific vitamin D receptor (VDR)-mediated transcriptional regulation of CYP3A4 gene. *Biochemical pharmacology* **79**, 277–287, doi: 10.1016/j.bcp.2009.08.017 (2010).
67. Lee, T. H. & Pelletier, J. Functional characterization of WT1 binding sites within the human vitamin D receptor gene promoter. *Physiological genomics* **7**, 187–200, doi: 10.1152/physiolgenomics.00046.2001 (2001).
68. Cockrell, A. S., Ma, H., Fu, K., McCown, T. J. & Kafri, T. A trans-lentiviral packaging cell line for high-titer conditional self-inactivating HIV-1 vectors. *Molecular therapy: the journal of the American Society of Gene Therapy* **14**, 276–284, doi: 10.1016/j.ymthe.2005.12.015 (2006).
69. Edgar, R., Domrachev, M. & Lash, A. E. Gene Expression Omnibus: NCBI gene expression and hybridization array data repository. *Nucleic acids research* **30**, 207–210 (2002).
70. Prat, A. *et al.* Phenotypic and molecular characterization of the claudin-low intrinsic subtype of breast cancer. *Breast cancer research: BCR* **12**, R68, doi: 10.1186/bcr2635 (2010).

Acknowledgements

We thank all the members of the Hohenstein and Hastie labs for the helpful discussions, Matthew Pearson for the imaging, Elisabeth Freyer for the flow cytometry, Tatjana Sauka-Spengler and Ahmed A. Ahmed for their help and support throughout the revision process. AHS is very grateful for funding from Breast Cancer Now. PH was supported by the MRC (grant MR/M010341/1). The Roslin Institute receives Institute Strategic Programme grant funding from the Biotechnology and Biological Sciences Research Council (BBSRC grant BB/J004316/1).

Author Contributions

P.H., N.D.H. and M.A. conceived and designed this work; M.A., J.S., M.M. and A.T. performed the experiments; M.A., A.H.S., S.A., J.D.P. and L.R.M. analysed the data; E.K. provided the clinical samples; E.K. and V.B. discussed the experiments. M.A. wrote the paper, P.H. and A.H.S. edited it. All authors read and approved the final manuscript.

Additional Information

Supplementary information accompanies this paper at <http://www.nature.com/srep>

Competing Interests: The authors declare no competing financial interests.

How to cite this article: Artibani, M. *et al.* *WT1* expression in breast cancer disrupts the epithelial/mesenchymal balance of tumour cells and correlates with the metabolic response to docetaxel. *Sci. Rep.* 7, 45255; doi: 10.1038/srep45255 (2017).

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>

© The Author(s) 2017